## IN THE CLAIMS

Please amend the claims as follows:

Claim 1 (Original): A crystalline, hydrated form of the sodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid, wherein the form contains from 6.4 up to 22 weight % of sodium and 15 up to 23 weight % of crystalline water if the sodium content is lower than 7.5 weight %, based on the whole molecule, or 4.5 up to 18 weight % of crystalline water if the sodium content is equal to or higher than 13 weight %, based on the anhydrous substance.

Claim 2 (Original): The crystalline form according to claim 1, which is pentahydrate of the monosodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid, wherein said form contains 20 up to 23 weight % of water built in the crystal lattice and 5.5 up to 7.5 % of sodium, based on the whole molecule.

Claim 3 (Currently Amended): The crystalline form according to claim 2, wherein said form eentains comprises 22.8 weight % of water built in the crystal lattice and 6.4 up to 6.7 % of sodium, based on the whole molecule.

Claim 4 (Currently Amended): The crystalline form according to claims 2 or 3, wherein said form shows a powder X-ray diffraction pattern with interplanar distances d that are approximately 16.3; 13.0; 9.1 and 4.9 Å.

Claim 5 (Currently Amended): The crystalline form according to claims 2 or 3, wherein said form shows the an infrared spectrum with having bands at 1169; 1060; 1046 and 891 cm<sup>-1</sup>.

Claim 6 (Currently Amended): The crystalline form according to claims 2 or 3, wherein a thermogravimetric analysis of which said form shows a plateau at temperature of about 173 °C.

Claim 7 (Currently Amended): The crystalline form according to claims 2 or 3, wherein a the <sup>31</sup>P CP-MAS NMR spectrum of which said form shows signals 13.7 and 20.0 ppm.

Claim 8 (Currently Amended): The crystalline form according to claim 1, which is trihydrate of the trisodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid, wherein said form contains comprises 19 up to 21 weight % of sodium and 12 up to 14 weight % of water in the crystal lattice.

Claim 9 (Currently Amended): The crystalline form according to claim 8, wherein said form shows the <u>an</u> infrared spectrum with <u>having</u> bands approximately <u>at</u> 1114; 1085; 956; 616 and 544 cm<sup>-1</sup>.

Claim 10 (Currently Amended): The crystalline form according to claim 1, which is monohydrate of the disodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid, wherein said form contains comprises 13 up to 15 weight % of sodium, based on the anhydrous substance, and 4.5 up to 6.5 weight % of water in the crystal lattice.

Claim 11 (Currently Amended): The crystalline form according to claim 10, wherein said form shows the an infrared spectrum with having bands at approximately 1183; 1158; 1071 and 1042 cm<sup>-1</sup>.

Claim 12 (Currently Amended): A method of manufacture of the crystalline form according to any of the foregoing claims characterized in that claim 1, comprising incorporating an aqueous solution of the sodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid heated to 50 up to 80 °C is incorporated into an organic solvent.

Claim 13 (Currently Amended): The method according to claim 12, wherein characterized in that the organic solvent is selected from the group consisting of simple alcohols from the C<sub>1</sub> to C<sub>5</sub> series, especially 2 propanol.

Claim 14 (Currently Amended): A method of manufacture of the crystalline form according to any of claims 1 through 11 characterized in that claim 1, comprising introducing seeding crystals of the respective hydrate of the sodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonate are introduced into the solution of the sodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid and slowly cooling the solution is slowly cooled.

Claim 15 (Currently Amended): The method according to claim 14, wherein characterized in that crystallization is performed from a solution of the sodium salt in a mixture of water and a water-miscible organic substance.

Claim 16 (Currently Amended): A pharmaceutical composition designed for treating diseases associated with a bone-resorption disorder, comprising characterized in that it eontains as the active agent, a hydrate of the sodium salt of 3-pyridyl-1-hydroxyethylidene-1,1-bisphosphonic acid according to claim 1 any of claims 1 through 11 and at least one auxiliary substance.

Claim 17 (New): The method according to claim 12, wherein the organic solvent is 2-propanol.